

118TH CONGRESS
1ST SESSION

H. R. 5647

To amend title XXVII of the Public Health Service Act to require out-of-network coverage for qualified individuals diagnosed with a rare pediatric disease participating in approved clinical trials, and for other purposes.

IN THE HOUSE OF REPRESENTATIVES

SEPTEMBER 21, 2023

Mr. McCaul (for himself, Mr. BERA, and Mr. KELLY of Pennsylvania) introduced the following bill; which was referred to the Committee on Energy and Commerce, and in addition to the Committee on Ways and Means, for a period to be subsequently determined by the Speaker, in each case for consideration of such provisions as fall within the jurisdiction of the committee concerned

A BILL

To amend title XXVII of the Public Health Service Act to require out-of-network coverage for qualified individuals diagnosed with a rare pediatric disease participating in approved clinical trials, and for other purposes.

1 *Be it enacted by the Senate and House of Representa-
2 tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Childhood Cancer Clin-
5 ical Trials Act”.

1 **SEC. 2. AMENDMENTS RELATING TO COVERAGE IN INDIVIDUAL AND GROUP MARKET AND UNDER MEDICARE PROGRAM FOR QUALIFIED INDIVIDUALS DIAGNOSED WITH A RARE PEDIATRIC DISEASE PARTICIPATING IN APPROVED CLINICAL TRIALS.**

7 (a) INDIVIDUAL AND GROUP MARKET.—

8 (1) REQUIRING OUT-OF-NETWORK COVERAGE
9 OF ROUTINE PATIENT COSTS.—Section 2709 of the
10 Public Health Service Act (42 U.S.C. 300gg–8) is
11 amended—

12 (A) in subsection (a)(2)—

13 (i) in subparagraph (A), by inserting
14 “and subsection (c)(2)” after “(1)(B);
15 and

16 (ii) in subparagraph (B), by inserting
17 “and subsection (c)(2)” after “(1)(B);

18 (B) by amending subsection (c) to read as
19 follows:

20 “(c) TREATMENT OF NON-PARTICIPATING PROVIDERS.—

22 “(1) IN GENERAL.—Subject to paragraph (2),
23 this section shall not be construed to require a group
24 health plan, or a health insurance issuer offering
25 group or individual health insurance coverage, to
26 provide benefits for routine patient care services pro-

1 vided outside of the plan's (or coverage's) health
2 care provider network unless out-of-network benefits
3 are otherwise provided under the plan (or coverage).

4 “(2) QUALIFIED INDIVIDUALS DIAGNOSED
5 WITH A RARE PEDIATRIC DISEASE.—In the case of
6 a qualified individual diagnosed with a rare pediatric
7 disease, with respect to routine patient costs for
8 items and services furnished to such individual in
9 connection with participation in an approved clinical
10 trial by a nonparticipating provider, a group health
11 plan or health insurance issuer offering group or in-
12 dividual health insurance coverage that provides cov-
13 erage to such individual—

14 “(A) shall impose the same cost-sharing
15 requirement (expressed as a copayment amount
16 or coinsurance rate) that would apply if such
17 item or service was furnished by a participating
18 provider; and

19 “(B) shall pay to such nonparticipating
20 provider the amount by which the recognized
21 amount for such item or service exceeds the
22 cost-sharing amount for such item or service
23 (as determined in accordance with subpara-
24 graph (A)).”; and

(C) by adding at the end the following new subsection:

3 "(i) OTHER DEFINITIONS.—

4 “(1) NONPARTICIPATING PROVIDER; PARTICI-
5 PATING PROVIDER; RECOGNIZED AMOUNT.—For
6 purposes of this section, the terms ‘nonparticipating
7 provider’, ‘participating provider’, and ‘recognized
8 amount’ have the meaning given such terms in sec-
9 tion 2799A–1(a)(3).

10 “(2) RARE PEDIATRIC DISEASE.—The term
11 ‘rare pediatric disease’ has the meaning given such
12 term in section 529(a)(3) of the Federal Food,
13 Drug, and Cosmetic Act.”.

18 (A) by striking “include all items and serv-
19 ices” and inserting “include—

“(j) all items and services”; and

21 (B) by striking the period at the end and
22 inserting “: and

1 (3) AMENDMENT RELATING TO DEFINITION OF
2 APPROVED CLINICAL TRIAL.—Section 2709(d)(1)(A)
3 of the Public Health Service Act (42 U.S.C. 300gg–
4 8(d)(1)(A)) is amended by adding at the end the fol-
5 lowing new clause:

6 “(viii) The Patient-Centered Out-
7 comes Research Institute.”.

8 (4) TECHNICAL AND CONFORMING AMEND-
9 MENTS.—Section 2709 of the Public Health Service
10 Act (42 U.S.C. 300gg–8), as amended by the pre-
11 ceding paragraphs, is further amended—

12 (A) in subsection (a)—

13 (i) in paragraph (1)(A), by inserting
14 before “clinical trial referred to in sub-
15 section (b)(2)” the following: “approved”;

16 (ii) in paragraph (2)(A), by striking
17 “a clinical trial” and inserting “an ap-
18 proved clinical trial”;

19 (iii) in paragraph (3)—

20 (I) by striking “IN-NETWORK
21 PROVIDERS” and inserting “PARTICI-
22 PATING PROVIDERS”; and

23 (II) by striking “a clinical trial”
24 and inserting “an approved clinical
25 trial”; and

1 (iv) in paragraph (4), by striking
2 “OUT-OF-NETWORK” and inserting “NON-
3 PARTICIPATING PROVIDERS”;

(B) in subsection (b)(2)(A), by striking “participating health care provider” and inserting “participating provider”; and

10 (5) EFFECTIVE DATE.—The amendments made
11 by this subsection shall apply with respect to plan
12 years beginning on or after January 1, 2024.

13 (b) MEDICARE.—

21 (B) by adding at the end the following new
22 paragraph:

“(3) ROUTINE COSTS OF CARE.—In defining ‘routine costs of care’ for purposes of paragraph (1), the Secretary shall define such term in a manner

1 that provides for coverage of consultation and refer-
2 ral services furnished to an individual in the course
3 of participation in a category A clinical trial.”.

4 (2) AMENDMENT RELATING TO DEFINITION OF
5 CATEGORY A CLINICAL TRIAL.—Section 1862(m)(2)
6 of the Social Security Act (42 U.S.C. 1395y(m)(2))
7 is amended by inserting after “means a trial” the
8 following: “(including a trial funded by the Patient-
9 Centered Outcomes Research Institute)”.

10 (3) EFFECTIVE DATE.—The amendments made
11 by this subsection shall apply with respect to items
12 and services furnished on or after January 1, 2024.

13 **SEC. 3. VOLUNTARY NETWORK OF PARTICIPATING PRO-**
14 **VIDERS.**

15 (a) IN GENERAL.—The Secretary of Health and
16 Human Services may issue a request for information from
17 group health plans, and health insurance issuers offering
18 group or individual health coverage to identify an interest
19 in establishing a voluntary network of participating pro-
20 viders administered by a third-party administrator (as
21 designated by the Secretary) for purposes of complying
22 with coverage requirements for clinical trials under section
23 2709 of the Public Health Service Act (42 U.S.C. 300gg–
24 8).

25 (b) DEFINITIONS.—In this section:

1 (1) GROUP HEALTH PLAN.—The term “group
2 health plan” has the meaning given such term in
3 section 607(1) of the Employee Retirement Income
4 Security Act of 1974 (29 U.S.C. 1167(1)).

5 (2) HEALTH INSURANCE ISSUER.—The term
6 “health insurance issuer” has the meaning given
7 such term in section 2791(b)(1) of the Public Health
8 Service Act (42 U.S.C. 300gg–91(b)(1)).

9 (3) PARTICIPATING PROVIDER.—The term
10 “participating provider” has the meaning given such
11 term in section 2799A–1(a)(3)(G)(ii) of the Public
12 Health Service Act (42 U.S.C. 300gg–
13 111(a)(3)(G)(ii)).

